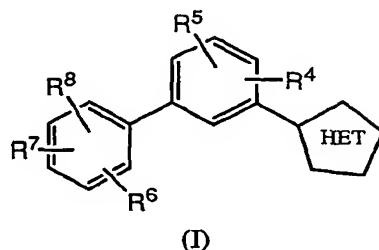


5 WHAT IS CLAIMED IS:

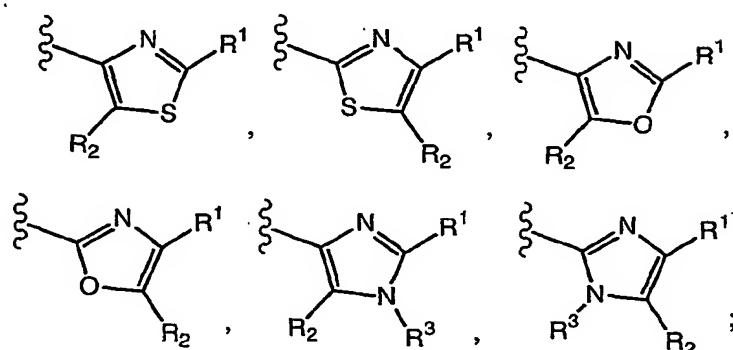
1. A compound represented by Formula (I):



10

or a pharmaceutically acceptable salt thereof, wherein

15 HET is one of the following heterocycles:



R1 is

(a) H;

20 (b) C₁-C₆-alkyl, C₂-C₄-alkenyl, C₂-C₄-alkynyl, C₃-C₆-cycloalkyl, or C₁-C₄-alkyl-[C₃-C₆-cycloalkyl], any of which is optionally substituted with one or more of the following substituents: F, CF₃, OH, O-(C₁-C₄)alkyl, S(O)₀₋₂-(C₁-C₄)alkyl, O-CONR^aR^b, NR^aR^b, N(R^a)CONR^aR^b, COO-(C₁-C₄)alkyl, COOH, CN, CONR^aR^b, SO₂NR^aR^b, N(R^a)SO₂NR^aR^b, -C(=NH)NH₂, tetrazolyl, triazolyl, imidazolyl, oxazolyl, oxadiazolyl, isooxazolyl, thiazolyl, furyl, thiienyl, pyrazolyl, pyrrolyl, pyridyl, pyrimidinyl, pyrazinyl, phenyl, piperidinyl, morpholinyl, pyrrolidinyl or piperazinyl;

25 (c) -O-C₁-C₆-alkyl, -O-C₃-C₆-cycloalkyl, -S-C₁-C₆-alkyl or -S-C₃-C₆-cycloalkyl, any of which is optionally substituted with one or more of the following substituents: F, CF₃, OH, O-(C₁-C₄)alkyl, S(O)₀₋₂-(C₁-C₄)alkyl, O-CONR^aR^b, NR^aR^b, N(R^a)CONR^aR^b, COO-(C₁-C₄)alkyl, COOH, CN, CONR^aR^b, SO₂NR^aR^b, N(R^a)SO₂NR^aR^b, -C(=NH)NH₂, tetrazolyl, triazolyl, imidazolyl, oxazolyl,

5 oxadiazolyl, isooxazolyl, thiazolyl, furyl, thienyl, pyrazolyl, pyrrolyl, pyridyl, pyrimidinyl, pyrazinyl, phenyl, piperidinyl, morpholinyl, pyrrolidinyl or piperazinyl;

(d) -C₀-C₄-alkyl-C₁-C₄-perfluoroalkyl, or -O-C₀-C₄-alkyl-C₁-C₄-perfluoroalkyl;

(e) -OH;

(f) -O-aryl, or -O-C₁-C₄-alkyl-aryl, wherein aryl is phenyl, pyridyl, pyrimidinyl, furyl, thienyl, pyrrolyl, triazolyl, pyrazolyl, thiazolyl, isoxazolyl, oxazolyl, or oxadiazolyl, any aryl of which is optionally substituted with 1-3 substituents selected from i) F, Cl, Br, I, ii) -CN, iii) -NO₂, iv) -C(=O)(R^a), v) -OR^a, vi) -NR^aR^b, vii) -C₀-4alkyl-CO-OR^a, viii) -(C₀-4alkyl)-NH-CO-OR^a, ix) -(C₀-4alkyl)-CO-N(R^a)(R^b), x) -S(O)₀₋₂R^a, xi) -SO₂N(R^a)(R^b), xii) -NR^aSO₂R^a, xiii) -C₁₋₁₀alkyl, and xiv) -C₁₋₁₀alkyl, wherein one or more of the alkyl carbons can be replaced by a -NR^a-, -O-, -S(O)₁₋₂-, -O-C(O)-, -C(O)-O-, -C(O)-N(R^a)-, -N(R^a)-C(O)-, -N(R^a)-C(O)-N(R^a)-, -C(O)-, -CH(OH)-, -C=C-, or -C≡C-;

(g) -OCON(R^a)(R^b), or -OSO₂N(R^a)(R^b);

(h) -SH, or -SCON(R^a)(R^b);

(i) NO₂;

20 (j) NR^aR^b, -N(COR^a)R^b, -N(SO₂R^a)R^b, -N(R^a)SO₂N(R^a)₂, -N(OR^a)CONR^aR^b, -N(R^a)SO₂R^a or -N(R^a)CON(R^a)₂;

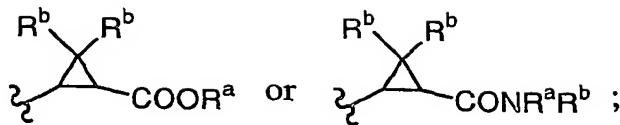
(k) -CH(OR^a)R^a, -C(OR^b)CF₃, -CH(NHR^b)R^a, -C(=O)R^a, C(=O)CF₃, -SOCH₃, -SO₂CH₃, COOR^a, CN, CONR^aR^b, -COCONR^aR^b, -SO₂NR^aR^b, -CH₂O-SO₂NR^aR^b, SO₂N(R^a)OR^a, -C(=NH)NH₂, -CR^a=N-OR^a, CH=CHCONR^aR^b;

25 (l) -CONR^a(CH₂)₀₋₂C(R^a)(R^b)(CH₂)₀₋₂CONR^aR^b;

(m) tetrazolyl, tetrazolinonyl, triazolyl, triazolinonyl, imidazolyl, imidozolonyl, oxazolyl, oxadiazolyl, isooxazolyl, thiazolyl, furyl, thienyl, pyrazolyl, pyrazolonyl, pyrrolyl, pyridyl, pyrimidinyl, pyrazinyl, or phenyl, any of which is optionally substituted with 1-3 substituents selected from i) F, Cl, Br, I, ii) -CN, iii) -NO₂, iv) -C(=O)R^a, v) C₁-C₆-alkyl, vi) -O-R^a, vii) -NR^aR^b, viii) -C₀-C₄-alkyl -CO-O R^a, ix) -(C₀-C₄-alkyl)-NH-CO-OR^a, x) -(C₀-C₄-alkyl)-CO-NR^aR^b, xi) -S(O)₀₋₂R^a, xii) -SO₂NR^aR^b, xiii) -NHSO₂R^a,xiv) -C₁-C₄-perfluoroalkyl, and xv) -O-C₁-C₄-perfluoroalkyl;

(n) -C(R^a)=C(R^b)-COOR^a, or -C(R^a)=C(R^b)-CONR^aR^b;

5 (o)



(p) piperidin-1-yl, morpholin-4-yl, pyrrolidin-1-yl, piperazin-1-yl or 4-susbtituted piperazin-1-yl, any of which is optionally substituted with 1-3 substituents selected from i) -CN, ii) -C(=O)(R^a), iii) C₁-C₆-alkyl, iv) -OR^a, v) -NR^aR^b, vi) -C₀-C₄-alkyl-CO-OR^a, vii) -(C₀-C₄-alkyl)-NH-CO-OR^a, viii) -(C₀-C₄-alkyl)-CON(R^a)(R^b), ix) -SR^a, x) -S(O)₀₋₂R^a, xi) -SO₂N(R^a)(R^b), xii) -NR^aSO₂R^a xiii) -C₁-C₄-perfluoroalkyl and xiv) -O-C₁-C₄-perfluoroalkyl;

R^a is

15 (a) H;

(b) C₁-C₄-alkyl, optionally substituted with one or more of the following substituents: F, CF₃, OH, O-(C₁-C₄)alkyl, S(O)₀₋₂(C₁-C₄)alkyl, -OCONH₂, -OCONH(C₁-C₄alkyl), -OCON(C₁-C₄alkyl)(C₁-C₄alkyl), -OCONHC₁-C₄alkyl-aryl, -OCON(C₁-C₄alkyl)(C₁-C₄alkyl-aryl), NH₂, NH(C₁-C₄alkyl), N(C₁-C₄alkyl)(C₁-C₄alkyl), NH(C₁-C₄alkyl-aryl), N(C₁-C₄alkyl)(C₁-C₄alkyl-aryl), NHCONH₂, NHCONH(C₁-C₄alkyl), NHCONH(C₁-C₄alkyl-aryl), -NHCON(C₁-C₄alkyl)(C₁-C₄alkyl), NHCON(C₁-C₄alkyl)(C₁-C₄alkyl-aryl), N(C₁-C₄alkyl)CON(C₁-C₄alkyl)(C₁-C₄alkyl), N(C₁-C₄alkyl)CON(C₁-C₄alkyl-aryl), COO-(C₁-C₄-alkyl), COOH, CN, CONH₂, CONH(C₁-C₄alkyl), CON(C₁-C₄alkyl)(C₁-C₄alkyl), SO₂NH₂, SO₂NH(C₁-C₄alkyl), SO₂NH(C₁-C₄alkyl-aryl), SO₂N(C₁-C₄alkyl)(C₁-C₄alkyl), NHSO₂NH₂, -C(=NH)NH₂, tetrazolyl, triazolyl, imidazolyl, oxazolyl, oxadiazolyl, 25 isooxazolyl, thiazolyl, furyl, thienyl, pyrazolyl, pyrrolyl, pyridyl, pyrimidinyl, pyrazinyl, phenyl, piperidinyl, morpholinyl, pyrrolidinyl or piperazinyl;

(c) C₀-C₄-alkyl-(C₁-C₄)-perfluoroalkyl; or

(d) C₁-C₄-alkyl-aryl, wherein aryl is phenyl, pyridyl, pyrimidinyl, furyl, thienyl, pyrrolyl, triazolyl, pyrazolyl, thiazolyl, isoxazolyl, oxazolyl, or oxadiazolyl, any aryl of which is optionally substituted with 1-3 substituents selected from i) F, Cl, Br, I, ii) -CN, iii) -NO₂, iv) -C(=O)(C₁-C₄-alkyl), v) -O(C₁-C₄-alkyl), vi) -N(C₁-C₄-alkyl)(C₁-C₄-alkyl), vii) -C₁-C₁₀alkyl, and viii) -C₁-C₁₀alkyl, wherein one or more of the alkyl carbons can be replaced by a -O-, -S(O)₁₋₂-, -O-C(O)-, -C(O)-O-, -C(O)-, -CH(OH)-, -C=C-, or -C≡C-;

35 R^b is

(a) H; or

5 (b) C_1-C_6 -alkyl, optionally substituted with one or more of the following substituents: F, CF_3 , OH, O-(C_1-C_4)alkyl, S(O)₀₋₂-(C_1-C_4)alkyl, -OCONH₂, -OCONH(C_1-C_4 alkyl), NH₂, NH(C_1-C_4 alkyl), N(C_1-C_4 alkyl)(C_1-C_4 alkyl), NHCONH₂, NHCONH(C_1-C_4 alkyl), -NHCON(C_1-C_4 alkyl)(C_1-C_4 alkyl), COO-(C_1-C_4 -alkyl), COOH, CN, and CONH₂;

10 R² is:

- (a) H;
- (b) - C_1-C_4 -alkyl, - C_3-C_6 -cycloalkyl or - C_1-C_4 -alkyl-(C_3-C_6)-cycloalkyl, optionally substituted with one or more of the following substituents: F, CF_3 , OH, O-(C_1-C_4)alkyl, S(O)₀₋₂-(C_1-C_4)alkyl, O-CONR^aR^b, NR^aR^b, N(R^a)CONR^aR^b, COO-(C_1-C_4)alkyl, COOH, CN, CONR^aR^b, SO₂NR^aR^b, N(R^a)SO₂NR^aR^b, -C(=NH)NH₂, tetrazolyl, triazolyl, imidazolyl, oxazolyl, oxadiazolyl, isooxazolyl, thiazolyl, furyl, thienyl, pyrazolyl, pyrrolyl, pyridyl, pyrimidinyl, pyrazinyl, phenyl, piperidinyl, morpholinyl, pyrrolidinyl and piperazinyl;
- (c) - C_0-C_4 -alkyl- C_1-C_4 -perfluoroalkyl;
- (d) aryl or -(C_1-C_4 -alkyl)-aryl, wherein aryl is phenyl, pyridyl, pyrimidinyl, furyl, thienyl, pyrrolyl, triazolyl, pyrazolyl, thiazolyl, isoxazolyl, oxazolyl, or oxadiazolyl, any aryl of which is optionally substituted with 1-3 substituents selected from i) F, Cl, Br, I, ii) -CN, iii) -NO₂, iv) -C(=O)(R^a), v) -OR^a, vi) -NR^aR^b, vii) -C₀₋₄alkyl-CO-OR^a, viii) -(C₀₋₄alkyl)-NH-CO-OR^a, ix) -(C₀₋₄alkyl)-CO-N(R^a)(R^b), x) -S(O)₀₋₂R^a, xi) -SO₂N(R^a)(R^b), xii) -NR^aSO₂R^a, xiii) -C₁₋₁₀alkyl, and xiv) -C₁₋₁₀alkyl, wherein one or more of the alkyl carbons can be replaced by a -NR^a-, -O-, -S(O)₁₋₂-, -O-C(O)-, -C(O)-O-, -C(O)-N(R^a)-, -N(R^a)-C(O)-, -N(R^a)-C(O)-N(R^a)-, -C(O)-, -CH(OH)-, -C=C-, or -C≡C-;
- (e) -C(=O)(R^a), -CONR^aR^b, COO-(C_1-C_4)alkyl, -SO₂R^a, -SO₂N(R^a)(R^b);

R³ is

- (a) H;
- (b) - C_1-C_4 -alkyl, - C_3-C_6 -cycloalkyl or - C_1-C_4 -alkyl-(C_3-C_6)-cycloalkyl, optionally substituted with one or more of the following substituents: F, CF_3 , OH, O-(C_1-C_4)alkyl, S(O)₀₋₂-(C_1-C_4)alkyl, O-CONR^aR^b, NR^aR^b, N(R^aR^b)CONR^aR^b, COO-(C_1-C_4)alkyl, COOH, CN, CONR^aR^b, SO₂NR^aR^b, N(R^aR^b)SO₂NR^aR^b, -C(=NH)NH₂, tetrazolyl, triazolyl, imidazolyl, oxazolyl, oxadiazolyl, isooxazolyl, thiazolyl, furyl, thienyl, pyrazolyl, pyrrolyl, pyridyl, pyrimidinyl, pyrazinyl, phenyl, piperidinyl, morpholinyl, pyrrolidinyl or piperazinyl;
- (c) - C_0-C_4 -alkyl- C_1-C_4 -perfluoroalkyl;
- (d) aryl or -(C_1-C_4 -alkyl)-aryl, wherein aryl is phenyl, pyridyl, pyrimidinyl, furyl, thienyl, pyrrolyl, triazolyl, pyrazolyl, thiazolyl, isoxazolyl, oxazolyl, or oxadiazolyl, any aryl of which is optionally substituted with 1-3 substituents selected from i) F, Cl, Br, I, ii) -CN, iii) -NO₂, iv) -C(=O)(R^a), v)

5 -OR^a, vi) -NR^aR^b, vii) -C₀₋₄alkyl-CO-OR^a, viii) -(C₀₋₄alkyl)-NH-CO-OR^a, ix) -(C₀₋₄alkyl)-CO-N(R^a)(R^b), x) -S(O)₀₋₂R^a, xi) -SO₂N(R^a)(R^b), xii) -NR^aSO₂R^a, xiii) -C₁₋₁₀alkyl, and xiv) -C₁₋₁₀alkyl, wherein one or more of the alkyl carbons can be replaced by a -NR^a- , -O-, -S(O)₁₋₂-, -O-C(O)-, -C(O)-O-, -C(O)-N(R^a)-, -N(R^a)-C(O)-, -N(R^a)-C(O)-N(R^a)-, -C(O)-, -CH(OH)-, -C=C-, or -C≡C- ;

(e) -O-C₁-C₄-alkyl, -O-C₀-C₄-alkyl-C₁-C₄-perfluoroalkyl, -O-aryl or -O(C₁-C₄-alkyl)-aryl;

10 (f) -C(=O)(R^a), -SO₂R^a, -SO₂N(R^a)(R^b), CN, NR^aR^b, NO₂, F, Cl, Br, I, OH, OCONR^aR^b, O(C₁-C₄-alkyl)CONR^aR^b, -OSO₂NR^aR^b, COOR^a, or CONR^aR^b;

R⁴ and R⁵ each independently is:

(a) H;

15 (b) -C₁-C₆-alkyl, -C₂-C₆-alkenyl, -C₂-C₆-alkynyl or -C₃-C₆-cycloalkyl, any of which is optionally substituted with one or more of the following substituents: F, CF₃, -O-(C₁-C₄)alkyl, CN, -N(R^a)(R^b), -N(R^a)CO-(C₁-C₄)alkyl, COOR^b, CON(R^a)(R^b) or phenyl;

(c) -O-C₀-C₆-alkyl, -O-aryl, or -O-C₁-C₄-alkyl-aryl, wherein aryl is phenyl, pyridyl, pyrimidinyl, furyl, thienyl, pyrrolyl, triazolyl, pyrazolyl, thiazolyl, isoxazolyl, oxazolyl, or oxadiazolyl, any aryl of which is optionally substituted with 1-3 substituents selected from i) F, Cl, Br, I, ii) -CN, iii) -NO₂, iv) -C(=O)(R^a), v) -OR^a, vi) -NR^aR^b, vii) -C₀₋₄alkyl-CO-OR^a, viii) -(C₀₋₄alkyl)-NH-CO-OR^a, ix) -(C₀₋₄alkyl)-CO-N(R^a)(R^b), x) -S(O)₀₋₂R^a, xi) -SO₂N(R^a)(R^b), xii) -NR^aSO₂R^a, xiii) -C₁₋₁₀alkyl, and xiv) -C₁₋₁₀alkyl, wherein one or more of the alkyl carbons can be replaced by a -NR^a- , -O-, -S(O)₁₋₂-, -O-C(O)-, -C(O)-O-, -C(O)-N(R^a)-,

20 (d) -N(R^a)-C(O)-, -N(R^a)-C(O)-N(R^a)-, -C(O)-, -CH(OH)-, -C=C-, or -C≡C-;

(e) -C₀-C₄-alkyl-C₁-C₄-perfluoroalkyl, or -O-C₀-C₄-alkyl-C₁-C₄-perfluoroalkyl; or

(e) CN, NH₂, NO₂, F, Cl, Br, I, OH, OCON(R^a)(R^b) O(C₁-C₄-alkyl)CONR^aR^b, -OSO₂N(R^a)(R^b), COOR^b, CON(R^a)(R^b), or aryl, wherein aryl is phenyl, pyridyl, pyrimidinyl, furyl, thienyl, pyrrolyl, triazolyl, pyrazolyl, thiazolyl, isoxazolyl, oxazolyl, or oxadiazolyl, any aryl of which is optionally substituted with 1-3 substituents selected from i) F, Cl, Br, I, ii) -CN, iii) -NO₂, iv) -C(=O)(R^a), v) -OR^a, vi) -NR^aR^b, vii) -C₀₋₄alkyl-CO-OR^a, viii) -(C₀₋₄alkyl)-NH-CO-OR^a, ix) -(C₀₋₄alkyl)-CO-N(R^a)(R^b), x) -S(O)₀₋₂R^a, xi) -SO₂N(R^a)(R^b), xii) -NR^aSO₂R^a, xiii) -C₁₋₁₀alkyl, and xiv) -C₁₋₁₀alkyl, wherein one or more of the alkyl carbons can be replaced by a -NR^a- , -O-, -S(O)₁₋₂-, -O-C(O)-, -C(O)-O-, -C(O)-N(R^a)-, -N(R^a)-C(O)-, -N(R^a)-C(O)-N(R^a)-, -C(O)-, -CH(OH)-, -C=C-, or -C≡C-; and

35 R⁶, R⁷ and R⁸ each independently is:

(a) H;

(b) C₁-C₆-alkyl, C₂-C₆-alkenyl, C₃-C₄-alkynyl or C₃-C₆-cycloalkyl, any of which is optionally substituted with one or more of the following substituents: F, CF₃, OH, O-(C₁-C₄)alkyl, OCON(R^a)(R^b), NR^aR^b,

5 COOR^a, CN, CONR^aR^b, N(R^a)CONR^aR^b, N(R^a)SO₂NR^aR^b, SO₂NR^aR^b, S(O)₀₋₂(C₁-C₄-alkyl), -C(=NH)NH₂, tetrazolyl, triazolyl, imidazolyl, oxazolyl, oxadiazolyl, isooxazolyl, thiazolyl, furyl, thienyl, pyrazolyl, pyrrolyl, pyridyl, pyrimidinyl, pyrazinyl, phenyl, piperidinyl, morpholinyl, pyrrolidinyl, or piperazinyl;

10 (c) -O-C₁-C₆-alkyl, -O-C₃-C₆-cycloalkyl, -S-C₁-C₆-alkyl or -S-C₃-C₆-cycloalkyl, any of which is optionally substituted with one or more of the following substituents: F, CF₃, OH, O-(C₁-C₄)alkyl, NH₂, NH(C₁-C₄-alkyl), N(C₁-C₄-alkyl)₂, COOH, CN, CONH₂, CONH(C₁-C₄-alkyl), CONH(C₁-C₄-alkyl)₂, SO₂NH₂, SO₂NH(C₁-C₄-alkyl), tetrazolyl, triazolyl, imidazolyl, oxazolyl, oxadiazolyl, isooxazolyl, thiazolyl, furyl, thienyl, pyrazolyl, pyrrolyl, pyridyl, pyrimidinyl, pyrazinyl, phenyl, piperidinyl, morpholinyl, pyrrolidinyl, piperazinyl;

15 (d) -C₀-C₄-alkyl-C₁-C₄-perfluoroalkyl, or -O-C₀-C₄-alkyl-C₁-C₄-perfluoroalkyl;

20 (e) -O-aryl, or -O-C₁-C₄-alkyl-aryl, wherein aryl is phenyl, pyridyl, pyrimidinyl, furyl, thienyl, pyrrolyl, triazolyl, pyrazolyl, thiazolyl, isoxazolyl, oxazolyl, or oxadiazolyl, any aryl of which is optionally substituted with 1-3 substituents selected from i) F, Cl, Br, I, ii) -CN, iii) -NO₂, iv) -C(=O)(R^a), v) -OR^a, vi) -NR^aR^b, vii) -C₀₋₄alkyl-CO-OR^a, viii) -(C₀₋₄alkyl)-NH-CO-OR^a, ix) -(C₀₋₄alkyl)-CO-N(R^a)(R^b), x) -S(O)₀₋₂R^a, xi) -SO₂N(R^a)(R^b), xii) -NR^aSO₂R^a, xiii) -C₁₋₁₀alkyl, and xiv) -C₁₋₁₀alkyl, wherein one or more of the alkyl carbons can be replaced by a -NR^a-, -O-, -S(O)₁₋₂-, -O-C(O)-, -C(O)-O-, -C(O)-N(R^a)-, -N(R^a)-C(O)-, -N(R^a)-C(O)-N(R^a)-, -C(O)-, -CH(OH)-, -C=C-, or -C≡C; (f) CN, N(R^a)(R^b), NO₂, F, Cl, Br, I, -OR^a, -SR^a, -OCON(R^a)(R^b), -OSO₂N(R^a)(R^b), COOR^b, CON(R^a)(R^b), -N(R^a)CON(R^a)(R^b), -N(R^a)SO₂N(R^a)(R^b), -C(OR^b)R^a, -C(OR^a)CF₃, -C(NHR^a)CF₃, -C(=O)R^a, C(=O)CF₃, -SOCH₃, -SO₂CH₃, -NHSO₂(C₁₋₆-alkyl), -NHSO₂-aryl, SO₂N(R^a)(R^b), -CH₂OSO₂N(R^a)(R^b), SO₂N(R^b)-OR^a, -C(=NH)NH₂, -CR_a=N-OR_a, CH=CH or aryl, wherein aryl is phenyl, pyridyl, pyrimidinyl, furyl, thienyl, pyrrolyl, triazolyl, pyrazolyl, thiazolyl, isoxazolyl, oxazolyl, or oxadiazolyl, any aryl of which is optionally substituted with 1-3 substituents selected from i) F, Cl, Br, I, ii) -CN, iii) -NO₂, iv) -C(=O)(R^a), v) -OR^a, vi) -NR^aR^b, vii) -C₀₋₄alkyl-CO-OR^a, viii) -(C₀₋₄alkyl)-NH-CO-OR^a, ix) -(C₀₋₄alkyl)-CO-N(R^a)(R^b), x) -S(O)₀₋₂R^a, xi) -SO₂N(R^a)(R^b), xii) -NR^aSO₂R^a, xiii) -C₁₋₁₀alkyl, and xiv) -C₁₋₁₀alkyl, wherein one or more of the alkyl carbons can be replaced by a -NR^a-, -O-, -S(O)₁₋₂-, -O-C(O)-, -C(O)-O-, -C(O)-N(R^a)-, -N(R^a)-C(O)-, -N(R^a)-C(O)-N(R^a)-, -C(O)-, -CH(OH)-, -C=C-, or -C≡C; or when R⁶ and R⁷ are present on adjacent carbon atoms, R⁶ and R⁷, together with the benzene ring to which they are attached, may form a bicyclic aromatic ring selected from naphthyl, indolyl, quinolinyl, isoquinolinyl, quinoxalinyl, benzofuryl, benzothienyl, benzoxazolyl, benzothiazolyl, and benzimidazolyl, any aromatic ring of which is optionally substituted with 1-4 independent substituents selected from i) halogen, ii) -CN, iii) -NO₂, iv) -CHO, v) -O-C₁₋₄alkyl, vi) -N(C₀₋₄alkyl)(C₀₋₄alkyl), vii) -C₀₋₄alkyl-CO-O(C₀₋₄alkyl), viii) -(C₀₋₄alkyl)-NH-CO-O(C₀₋₄alkyl), ix)

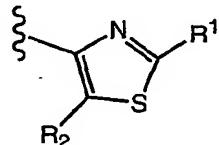
5 -(C₀₋₄alkyl)-CO-N(C₀₋₄alkyl)(C₀₋₄alkyl), x) -S(C₀₋₄alkyl), xi) -S(O)(C₁₋₄alkyl), xii) -SO₂(C₀₋₄alkyl), xiii) -SO₂N(C₀₋₄alkyl)(C₀₋₄alkyl), xiv) -NHSO₂(C₀₋₄alkyl)(C₀₋₄alkyl), xv) -C₁₋₁₀alkyl and xvi) -C₁₋₁₀alkyl in which one or more of the carbons can be replaced by a -N(C₀₋₆alkyl)-, -O-, -S(O)₁₋₂-, -O-C(O)-, -C(O)-O-, -C(O)-N(C₀₋₆alkyl)-, -N(C₀₋₆alkyl)-C(O)-, -N(C₀₋₆alkyl)-C(O)-N(C₀₋₆alkyl)-, -C(O)-, -CH(OH), -C=C-, or -C≡C-.

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2. A compound according to Claim 1, or a pharmaceutically acceptable salt thereof,
wherein

HET is

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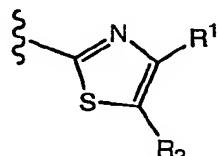


3. A compound according to Claim 1, or a pharmaceutically acceptable salt thereof,

wherein

20

HET is



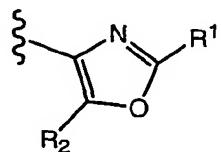
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4. A compound according to Claim 1, or a pharmaceutically acceptable salt thereof,

wherein

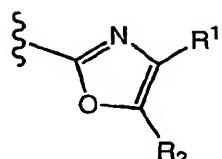
HET is

30



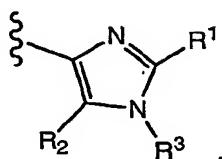
5 5. A compound according to Claim 1, or a pharmaceutically acceptable salt thereof,
wherein

HET is



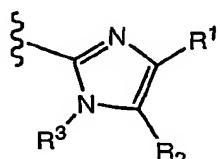
6. A compound according to Claim 1, or a pharmaceutically acceptable salt thereof,
wherein

15 HET is



20 7. A compound according to Claim 1, or a pharmaceutically acceptable salt thereof,
wherein

HET is

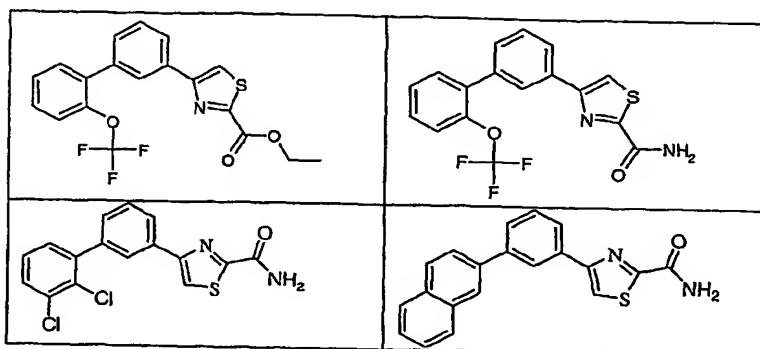


8. A compound according to Claim 1, or a pharmaceutically acceptable salt thereof,
wherein

30 R⁶ is other than H and is attached at the ortho position.

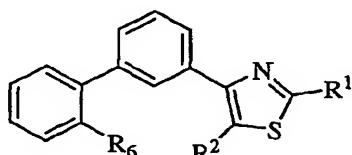
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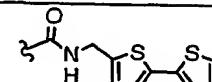
9. A compound represented by

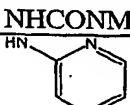
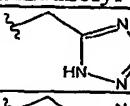
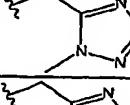


10. A compound according to Claim 1 represented by

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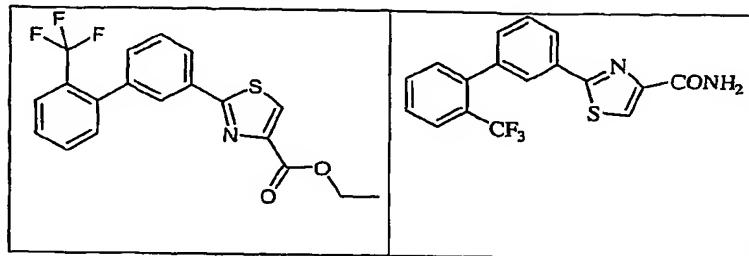


R⁶	R²	R¹
Cl	H	H
Cl	H	COOEt
Cl	H	CONH ₂
Cl	H	CONH-tBu
Cl	H	
Cl	H	NH ₂
CF ₃	H	COOEt
CF ₃	H	CONH ₂
CF ₃	H	H
CF ₃	H	NH ₂
OCF ₃	H	CH ₃
OCF ₃	H	H
OCF ₃	H	NH ₂
OCF ₃	H	CONMe ₂

R⁶	R²	R¹
OCF ₃	Cl	CH ₃
OCF ₃	H	NHSO ₂ CH ₃
OCF ₃	H	CH ₂ OH
O-Ph	H	CONH ₂
CF ₃	H	NHCONH-iPr
OCF ₃	H	NHCONH-iPr
OCF ₃	H	NHCOCH ₃
CF ₃	H	NHCOCH ₃
OCF ₃	H	CH ₂ COOEt
OCF ₃	H	CH ₂ CN
OCF ₃	H	CH ₂ CONH ₂
CF ₃	H	CH ₂ CONH ₂
OCF ₃	H	NHCONMe ₂
OCF ₃	H	
OCF ₃	H	2-Pyrimidyl
OCF ₃	H	2-Pyridyl
OCF ₃	H	2-Oxazolyl
OCF ₃	H	2-Imidazolyl
OCF ₃	H	2-Pyrazolyl
OCF ₃	H	2-(1-Methyl)-imidazolyl
OCF ₃	H	
OCF ₃	H	
OCF ₃	H	

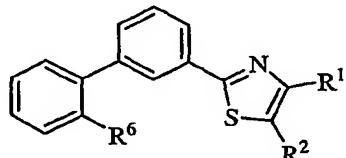
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11. A compound represented by



12. A compound according to Claim 1 represented by

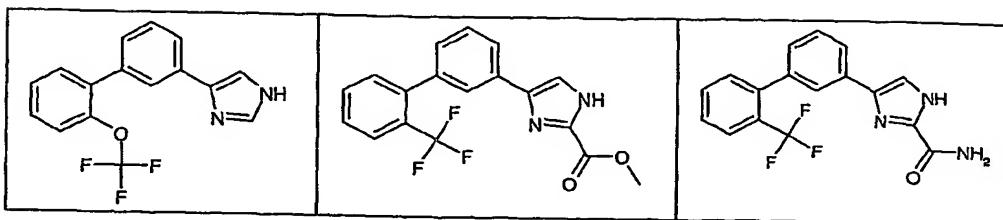
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R ₆	R ₂	R ₁
CF ₃	H	H
CF ₃	H	COOEt
CF ₃	H	CONH ₂
CF ₃	H	CONHCH ₃
CF ₃	COOEt	CH ₃
CF ₃	CONH ₂	CH ₃
OCF ₃	H	H
OCF ₃	H	COOCH ₃
OCF ₃	H	CONH ₂
OCF ₃	H	COOH
OCF ₃	H	CH ₂ OH
OCF ₃	H	CONH(CH ₂) ₃ OH
O-Ph	H	CONH ₂

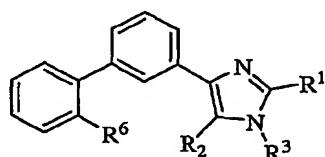
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13. A compound represented by



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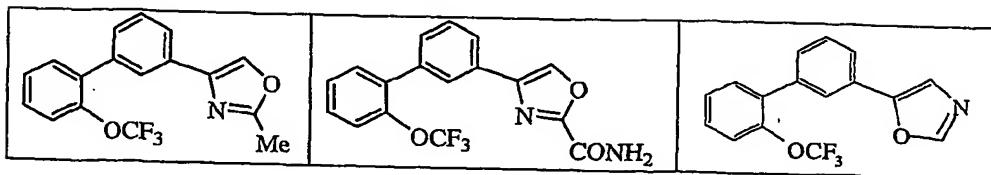
14. A compound according to Claim 1 represented by



R⁶	R³	R²	R¹
Cl	H	H	H
Cl	H	H	COOCH ₃
Cl	H	H	CONH ₂
CF ₃	H	H	H
CF ₃	H	H	COOH
CF ₃	CH ₃	H	CONH ₂
CF ₃	H	Cl	COOCH ₃
CF ₃	H	Cl	CONH ₂
CF ₃	H	Cl	Cl
OCF ₃	H	H	H
OCF ₃	H	H	COOCH ₃
OCF ₃	H	H	CONH ₂
OCF ₃	H	H	

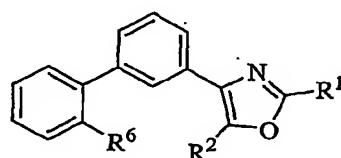
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15. A compound represented by



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16. A compound according to Claim 1 represented by



R ⁶	R ²	R ¹
Cl	H	CH ₃
Cl	H	NH ₂
CF ₃	H	CH ₃
OCF ₃	H	NH ₂
OCF ₃	H	CH=CH ₂
OCF ₃	H	COOCH ₃
OCF ₃	H	CONH ₂
OCF ₃	H	CH ₃
CF ₃	H	NH ₂
CF ₃	H	H
OCF ₃	H	H
Cl	H	COOH
CF ₃	H	CONH ₂
OCF ₃	H	COOCH ₂ CH ₃

15

17. A pharmaceutical composition comprising a therapeutically effective amount of the compound according to Claim 1, or a pharmaceutically acceptable salt thereof, and a pharmaceutically acceptable carrier.

5

18. The pharmaceutical composition according to Claim 17, further comprising a second therapeutic agent selected from the group consisting of: i) opiate agonists, ii) opiate antagonists, iii) calcium channel antagonists, iv) 5HT receptor agonists, v) 5HT receptor antagonists vi) sodium channel antagonists, vii) NMDA receptor agonists, viii) NMDA receptor antagonists, ix) COX-2 selective inhibitors, x) NK1 antagonists, xi) non-steroidal anti-inflammatory drugs, xii) selective serotonin reuptake inhibitors, xiii) selective serotonin and norepinephrine reuptake inhibitors, xiv) tricyclic antidepressant drugs, xv) norepinephrine modulators, xvi) lithium, xvii) valproate xviii) neurontin, and xix) antiviral agents.

10

15 19. A method of treatment or prevention of pain comprising the step of administering to a patient in need thereof a therapeutically effective amount, or a prophylactically effective amount, of the compound according to Claim 1, or a pharmaceutically acceptable salt thereof.

20

20. A method of treatment of chronic, visceral, inflammatory and neuropathic pain syndromes comprising the step of administering to a patient in need thereof a therapeutically effective amount, or a prophylactically effective amount, of the compound according to Claim 1, or a pharmaceutically acceptable salt thereof.

25

21. A method of treatment of pain resulting from, or associated with, traumatic nerve injury, nerve compression or entrapment, postherpetic neuralgia, trigeminal neuralgia, diabetic neuropathy, cancer and chemotherapy, comprising the step of administering to a patient in need thereof a therapeutically effective amount, or a prophylactically effective amount, of the compound according to Claim 1, or a pharmaceutically acceptable salt thereof.

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22. A method of treatment of chronic lower back pain comprising the step of administering to a patient in need thereof a therapeutically effective amount, or a prophylactically effective amount, of the compound according to Claim 1, or a pharmaceutically acceptable salt thereof.

35

23. A method of treatment of phantom limb pain comprising the step of administering to a patient in need thereof a therapeutically effective amount, or a prophylactically effective amount, of the compound according to Claim 1, or a pharmaceutically acceptable salt thereof.

24. A method of treatment of HIV- and HTV treatment-induced neuropathy, chronic pelvic pain, neuroma pain, complex regional pain syndrome, chronic arthritic pain and related neuralgias

5 comprising the step of administering to a patient in need thereof a therapeutically effective amount, or a prophylactically effective amount, of the compound according to Claim 1, or a pharmaceutically acceptable salt thereof.

10 25. A method of administering local anesthesia comprising the step of administering to a patient in need thereof a therapeutically effective amount, or a prophylactically effective amount, of the compound according to Claim 1, or a pharmaceutically acceptable salt thereof.

15 26. A method of treatment of irritable bowel syndrome and Crohn's disease comprising the step of administering to a patient in need thereof a therapeutically effective amount, or a prophylactically effective amount, of the compound according to Claim 1, or a pharmaceutically acceptable salt thereof.

20 27. A method of treatment of epilepsy and partial and generalized tonic seizures comprising the step of administering to a patient in need thereof a therapeutically effective amount, or a prophylactically effective amount, of the compound according to Claim 1, or a pharmaceutically acceptable salt thereof.

25 28. A method for neuroprotection under ischaemic conditions caused by stroke or neural trauma comprising the step of administering to a patient in need thereof a therapeutically effective amount, or a prophylactically effective amount, of the compound according to Claim 1, or a pharmaceutically acceptable salt thereof.

30 29. A method of treatment of multiple sclerosis comprising the step of administering to a patient in need thereof a therapeutically effective amount, or a prophylactically effective amount, of the compound according to Claim 1, or a pharmaceutically acceptable salt thereof.

35 30. A method of treatment of bipolar disorder comprising the step of administering to a patient in need thereof a therapeutically effective amount, or a prophylactically effective amount, of the compound according to Claim 1, or a pharmaceutically acceptable salt thereof.

31. A method of treatment of tachy-arrhythmias comprising the step of administering to a patient in need thereof a therapeutically effective amount, or a prophylactically effective amount, of the compound according to Claim 1, or a pharmaceutically acceptable salt thereof.